EVIDENCE FOR NEW SULFATE PARTICLE FORMATION IN THE REMOTE TROPOSPHERE INVOLVING BIOGENIC TRACE GAS SPECIES

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Evidence For New Sulfate Particle Formation In The Remote Troposphere Involving Biogenic Trace Gas Species

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Findings from earlier field studies (Weber et al., 1995; Weber et al., 1997) showed that rates of new particle formation in the remote troposphere can be significantly higher than those predicted by classical heteromolecular sulfuric acid - water (H_2SO_4 - H_2O) nucleation theory (Jaecker-Voirol and Mirabel, 1989). We have speculated that higher rates may be due to the participation of ammonia (NH_3) through a ternary mechanism involving H_2SO_4 - NH_3 - H_2O (Weber et al., 1996). Airborne measurements made in the vicinity of Macquarie Island during the first Aerosol Characterization Experiment (ACE 1) support the hypothesis that participation of additional species can result in particle formation rates that significantly exceed rates for H_2SO_4 - H_2O nucleation (Weber et al., 1998a).

ACE 1 airborne measurements pertinent to studies of new particle formation included various meteorological parameters, gas phase H₂SO₄ and H₂O, and nanoparticle size distributions (~3 to 10 nm diameter). Nanoparticle concentrations were determined from the photo detector pulse heights of an Ultrafine Condensation Particle Counter (Weber et al., 1998b).

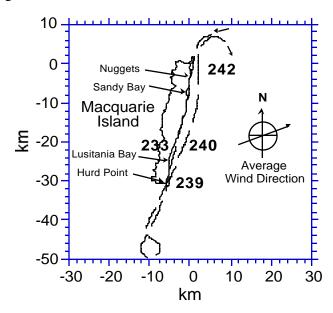


Figure 1: Aircraft measurements off the lee coast of Macquarie Island. Bands show aircraft trajectories during the one minute sample interval to measure nanoparticle spectra. Locations of four of the large penguin rookeries on Macquarie Island are identified by name. Thick bands, identified by sample number, are the only measurements where evidence of nucleation was observed. The corresponding nanoparticle distributions are shown in Figure 2.

Off the lee coast of Macquarie Island, measured nanoparticle size distributions showed clear evidence of nucleation exclusively downwind of large penguin rookeries (Figures 1 and 2), locations where NH₃ concentrations are expected to be high (Lindeboom, 1984). Of the measurements downwind of the rookeries, highest nanoparticle concentrations (e.g., sample number 240 in Figure 2) were observed in regions of highest H₂SO₄ concentrations, suggesting that H₂SO₄ also participated in particle formation. However, calculations show that in these regions nucleation of H₂SO₄ and H₂O would not occur.

This paper will present the Macquarie Island measurements and explore mechanisms of other nucleation events observed in the remote troposphere by comparing them to the Macquarie Island nucleation events and predictions of H₂SO₄-H₂O nucleation.

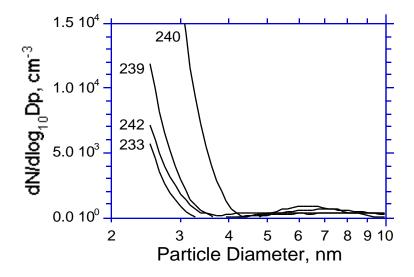


Figure 2: Nanoparticle size distributions recorded immediately downwind of penguin rookeries during ACE 1.

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